IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Patent application of: SCHUTTE, Brian C., et al.

For: IRF6 POLYMORPHISMS ASSOCIATED WITH CLEFT LIP AND/OR PALATE

the specification of which is being transmitted herewith.

INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

Attached are Forms PTO/SB/08B (formerly Form PTO-1449) listing the relevant art known to the applicant herein. Copies of the references are enclosed. The Examiner is requested to consider the references and make them of record.

Applicants disclose herewith patents, publications or other information, of which they are aware that they believe may be material to the examination of this application, and in respect of which, there may be a duty to disclose. Legible copies of all items listed in Forms PTO/SB/08B (formerly Form PTO-1449) accompany this information statement, except those identified above.

The filing of this information disclosure statement shall not be construed as a representation that a search has been made (37 C.F.R. § 1.97(g)), an admission that the

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information cited is, or is considered to be, material to patentability, or that no other material information exists.

The filing of this information disclosure statement shall not be construed as an admission against interest in any manner. (Notice of January 9, 1992, 1135 O.G. 13-25, at 25.)

Respectfully submitted,

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- bja -

Form PTO/SB/08B (10-01)
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Substitute for form 1449B/PTO				Complete if Known		
				Application Number		
INFO	RMATION D	ISCI	LOSURE	Filing Date	May 6, 2004	
	TEMENT BY			First Named Inventor	SCHUTTE, Brian C., et al.	
				Group Art Unit		
(Use as many sheets as necessary)				Examiner Name		
Sheet	1	of	1	Attorney Docket Number	P06215US01	

OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS						
Examiner Initials *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T 2			
	1	Kayano, et al., National Library of Medicine "Novel IRF6 mutations in Japanese patients with Van der Woude syndrome: to missense mutations (R45Q and P396S) and a 17-kb deletion." Abstract (1 page) 11/15/2003				
	2	Schutte, Brian C., et al. "A Preliminary Gene Map for the Van der Woude Syndrome Critical Region Derived from 900 kb of Geno,mic Sequence at 1q32-q41" Genome Research, 11/9/999				
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